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Maximilianstrasse 54
D-80538 München (DE)(54) **PROCESS FOR PRODUCING L-THREONINE BY FERMENTATION.**

(57) A DNA containing a gene that codes for aspartokinase III from a bacterium of the genus *Escherichia* and has in the coding region thereof a variation that releases the inhibition of aspartokinase III feedback by lysine; a recombinant DNA comprising the above DNA joined to a vector DNA capable of autonomous replication in the cell of a bacterium of the genus *Escherichia*; a micro-organism of the genus *Escherichia*, that has been transformed by introducing the recombinant DNA into the cells; and a process for producing L-threonine by culturing the micro-organism in a fermentation medium to accumulate L-threonine in the medium and harvesting the same from the medium. The invention process has provided AK III genes originating in a bacterium of the genus *Escherichia* and sufficiently released from the feedback inhibition caused by lysine. An improved threonine-producing bacterium has been obtained by introducing the above genes into a conventional threonine-producing bacterium. The invention successfully provides a process for producing L-threonine by an improved fermentation method using the above threonine-producing bacterium.

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(Technical Field)

The present invention relates to the microbiological industry, and describes a method for the production of L-threonine by fermentation. L-threonine is an essential amino acid, and it is used as a component of various nutritional mixtures for medical purposes. Furthermore, it is used as a feed additive for animals, as a reagent in the pharmaceutical and chemical industries, and as a growth factor for the production of amino acids such as lysine and homoserine by microorganisms.

(Background Art)

In the past, bacteria separated from the natural environment or artificial mutants of such bacteria have been used as microorganisms for the production of L-threonine by fermentation. Many L-threonine-producing artificial mutants are known, the majority of which are resistant to α -amino- β -hydroxyvaleric acid and belong to the genus *Escherichia*, *Serratia*, *Brevibacterium* or *Corynebacterium*. Regarding *Escherichia*, in JPA 55-131397, JPA 59-31691, JPA 56-15696 and JPW 3-501682 are there described methods for the production of L-threonine by bacteria transformed by recombinant plasmids containing threonine operons.

Also, of the heretofore known threonine-producing bacteria, *Escherichia coli* ВКПМ (VKPM) B-3996 strain has exhibited the most superior level of threonine production and coefficient of consumption (JPW 3-501682). According to the present invention, "coefficient of consumption" refers to the number of grams of sugar required for the production of one gram of threonine. If this bacterium is cultured in an experimental fermenter, adding sugar-ammonia to the culture medium in response to the signals from the pH sensor, then the maximum degree of biosynthesis of threonine is 85 g/l, and the coefficient of consumption is 2 g of sugar per one gram of threonine.

Aspartokinase (hereunder abbreviated to AK) is an enzyme which converts aspartic acid to β -phosphoaspartic acid, and it is the main regulatory site of the biosynthesis pathway of aspartic acid and its derivative amino acids. As shown in Fig. 1, there are three types of AK from *E. coli* (AK I, AK II, AK III) and the first two of these are bifunctional enzymes having homoserine dehydrogenase (hereunder sometimes abbreviated to HD) activity. One of these is AK I-HD I which is coded for by the thrA gene, and the other is AK II-HD II which is coded for the metL(M) gene.

Only AK III is a monofunctional enzyme, and it is the product of a gene named lysC, and is known to undergo repression and feedback inhibition by lysine. On the other hand, AK I undergoes concerted repression by threonine and isoleucine, and inhibition by threonine, while AK II undergoes repression by methionine. The proportion of the intracellular activities thereof is AK I:AK II:AK III = approximately 5:1:4.

The lysC gene of *E. coli* has already been cloned, and its base sequence has been determined (Cassan, M., Parsot, C., Cohen, G.N. and Patte, J.C., J. Biol. Chem., 261, 1052, 1986). Also, the production of L-threonine by fermentation using strains of *E. coli* whose AK III activity has been reinforced, is described by Mizukami, et al. (Mizukami, T. et al., Agric. Biol. Chem., 50, 1015, 1986). However, sufficient release of the lysine-dependent feedback inhibition on the AK III reported by Mizukami, et al. has not yet been achieved.

Thus, the subject matter of the present invention is obtaining AK III with sufficient release of the feedback inhibition due to lysine, and providing a method for the production of L-threonine by fermentation which is much improved over the prior art.

(Disclosure of Invention)

The inventors of the present invention, as a result of diligent research carried out to overcome the above mentioned problem, have succeeded in obtaining a gene (mutant lysC, or lysC*) which codes for AK III in *E. coli* and which sufficiently releases the feedback inhibition due to lysine, and have discovered that by introducing this gene into threonine-producing bacteria, L-threonine is efficiently accumulated, and thus the present invention has been completed.

In other words, the present invention relates to a DNA containing a gene which codes for aspartokinase III found in *Escherichia* bacteria and which has a mutation in the coding region which releases the feedback inhibition by lysine on the above mentioned aspartokinase III, and specifically, it relates to the above mentioned DNA in which the mutation which releases the feedback inhibition by lysine on aspartokinase III is positioned, e.g., on the A domain or B domain of aspartokinase III, and more specifically, it relates to the above mentioned DNA in which the mutation which releases the feedback inhibition by lysine on aspartokinase III is, with reference to the below mentioned Sequence Listing: SEQ ID NO: 1,

selected from the group consisting, e.g., of a mutation wherein the 323rd Gly is replaced by Asp; a mutation wherein the 323rd Gly is replaced by Asp and the 408th Gly is replaced by Asp; a mutation wherein the 34th Arg is replaced by Cys and the 323rd Gly is replaced by Asp; a mutation wherein the 325th Leu is replaced by Phe; a mutation wherein the 318th Met is replaced by Ile; a mutation wherein the 318th Met is replaced by Ile and the 349th Val is replaced by Met; a mutation wherein the 345th Ser is replaced by Leu; a mutation wherein the 347th Val is replaced by Met; a mutation wherein the 352nd Thr is replaced by Ile; a mutation wherein the 352nd Thr is replaced by Ile and the 369th Ser is replaced by Phe; a mutation wherein the 164th Glu is replaced by Lys; and a mutation wherein the 417th Met is replaced by Ile and the 419th Cys is replaced by Tyr.

Furthermore, the present invention relates to the above mentioned DNA which is a recombinant DNA linked with vector DNA capable of autonomous replication in *Escherichia* bacteria. Also, it relates to microorganisms belonging to the genus *Escherichia* which have been transformed by the introduction of the above mentioned recombinant DNA into the cells. Also within the scope of the present invention are microorganisms which have been transformed by the introduction into their chromosomal DNA of the DNA which codes for the above mentioned *Escherichia*-derived aspartokinase III and contains a gene with a mutation in the coding region which releases the feedback inhibition by lysine on the above mentioned aspartokinase III.

The present invention also relates to a method for the production of L-threonine, characterized by culturing any of the above mentioned microorganisms in a fermentation medium, producing and accumulating L-threonine in the culture, and collecting the L-threonine from the culture.

A detailed description of the present invention will now be provided.

As the donor bacteria to supply the DNA containing the gene coding for AK III (lysC) may be used any microorganism belonging to the genus *Escherichia*. Specifically, those mentioned in the writings of Neidhardt (Neidhardt, F.C. et al., *Escherichia coli* and *Salmonella typhimurium*, American Society for Microbiology, Washington, D.C., 1208, Table 1). Examples thereof include *E. coli* strains JM109 and MC1061.

If a wild strain is used as the donor bacteria to supply the DNA containing the gene coding for AK III (lysC), then a DNA containing a wild type of the AK III gene may be obtained. For the introduction of a mutation into this gene in order to obtain an AK III gene which releases the feedback inhibition due to L-lysine (lysC⁻), *in vitro* mutation of the DNA may be effected by direct treatment with hydroxylamine. Hydroxylamine is a chemical mutagenic agent which induces a mutation C→T by converting cytosine to N⁴-hydroxycytosine.

Further, a DNA containing the AK III gene which releases the feedback inhibition due to L-lysine may be obtained by using as a DNA donor strain a mutant which releases the feedback inhibition on AK III activity due to L-lysine. This mutant may be obtained from cells which have been subjected to, for example, a conventional method for mutagenic treatment, exposure to ultraviolet rays or treatment with a mutagenic agent such as N-methyl-N'-nitro-N-nitrosoguanidine (NTG).

A description will now be provided regarding the preparation of a DNA containing a gene which codes for AK III (lysC). First, *E. coli* having a wild type of lysC, for example, MC1061 strain, is cultured to obtain cultured cells. The culturing of the above mentioned microorganism may be carried out by a conventional solid culture method, but a liquid culture method is preferably used for the culturing from the view point of cell collection. Also, the culture medium may be one prepared for example, by adding one or more inorganic salts such as monopotassium phosphate (KH₂PO₃), dipotassium phosphate (K₂HPO₃), magnesium sulfate, sodium chloride, magnesium chloride, ferric chloride, ferric sulfate, or manganese sulfate, to one or more nitrogen sources such as yeast extract, peptone, beef extract, corn steep liquor or an effusion of soybean or wheat, and then further adding carbohydrates, vitamins, etc., if necessary. The initial pH of the culture medium is appropriately adjusted to 7-8. Also, the culturing is carried out at 30-42°C, and preferably about 37°C, for 4-24 hours, by submerged culture with aeration and stirring, shaking culture, standing culture, or the like. The cultured product which is obtained in this manner is subjected to centrifugal separation at, for example, 3,000 rpm for 5 minutes, to obtain cells of *E. coli* MC1061 strain.

Chromosomal DNA may be obtained from these cells by, for example, the method of Saito and Miura (Biochem. Biophys. Acta. 72, 619, 1963), the method of K.S. Kirby (Biochem. J. 64, 405, 1956), etc.

For isolating the lysC gene, the chromosomal DNA obtained by the above methods is cleaved using an appropriate restriction enzyme. If the degree of cleavage is controlled by controlling the cleavage reaction time, etc., then a wide variety of restriction enzymes may be used. Next, the gene may be linked to a vector DNA which is capable of replication in *Escherichia* bacteria, and the resulting recombinant DNA may be used to transform a mutant strain of *Escherichia*, for example, GT3, which lacks aspartokinase I, II and III (to create a gene library), and of the resulting transformants may be isolated a strain which has become

capable of growing on a minimal culture medium without lysine, thus separating the recombinant DNA containing the *lysC* gene.

Concretely, a chromosomal DNA is subjected to digestion with a restriction enzyme, for example, *Sau3AI* at a temperature of 30°C or higher, and preferably 37°C, at an enzyme concentration of 1-10 units/ml for various times (1 minute - 2 hours) for complete digestion or partial digestion to obtain a mixture containing a variety of chromosomal DNA fragments. The vector DNA which is capable of replication in *Escherichia* bacteria is subjected to digestion with a restriction enzyme, for example, *BamHI*, which produces the same terminal base sequence as does the restriction enzyme *Sau3A* used for the cleavage of the chromosomal DNA, at a temperature of 30°C or higher, at an enzyme concentration of 1-100 units/ml for 1 hour or more, and preferably 1-3 hours, to effect complete digestion thereof to obtain the cleaved DNA. Next, the mixture containing the DNA fragments which were derived from *E. coli* MC1061 and include the *lysC* gene prepared in the manner described above, is mixed with the cleaved vector DNA, and a DNA ligase, preferably T4 DNA ligase, is allowed to act thereon at a temperature of 4-16°C, at an enzyme concentration of 1-100 units/ml for 1 hour or more, preferably 6-24 hours, to obtain a recombinant DNA.

The vector DNA to be used according to the present invention is preferably a plasmid vector DNA, for example, pUC19, pUC18, pBR322, pHSG299, pHSG399, RSF1010, etc. In addition, phage DNA vectors may be used. Promoters which function in microorganisms, for example, *lac*, *trp*, *PL* and the like may be used for an efficient expression of the gene which is useful for the desired purpose. The term "recombinant DNA" used here includes DNA resulting from the incorporation of the above gene into chromosomes by methods which make use of transposons (Berg, D.E. and Berg, C.M., *Bio/Technol.*, 1, 417, 1983), Mu phage (Japanese Patent Application Disclosure HEI 2-109985) or homologous recombination (Experiments in Molecular Genetics, Cold Spring Harbor Lab., 1972).

This recombinant DNA is used to transform, for example, *E. coli* K-12 strain, and preferably GT3 strain, etc., and then a strain which has recombinant DNA containing the *lysC* gene is obtained from the strains having an increased level of AK activity or from strains with complemented nutritional requirements. The transformation may be carried out according to the method of D.M. Morrison (*Methods in Enzymology*, 68, 326, 1979) or a method whereby the recipient cells are treated with calcium chloride to raise the degree of penetration of the DNA (Mandel, M. and Higa, A., *J. Mol. Biol.*, 53, 159, 1970). Then, the recombinant DNA prepared by the insertion of DNA containing the *lysC* gene into the vector DNA may be isolated from the above mentioned strains following, for example, the method of P. Guerry, et al. (*J. Bacteriol.*, 116, 1064, 1973) or the method of D.B. Clewell (*J. Bacteriol.*, 110, 667, 1972).

It may be confirmed that the candidate strain in fact possesses the recombinant DNA having *lysC*, by preparing a cell extract solution, and then preparing therefrom a crude enzyme solution for confirmation of aspartokinase activity. The method of measuring the enzyme activity of aspartokinase may be according to the method of Stadtman, et al. (Stadtman, E.R., Cohen, G.N., LeBras, G., and Robichon-Szulmajster, H., *J. Biol. Chem.*, 236, 2033, 1961).

Alternatively, the *lysC* gene may be obtained by amplifying the *lysC* gene from the chromosomal DNA obtained by the method of Saito and Miura, by the PCR (polymerase chain reaction; see White, T.J. et al., *Trends Genet.* 5, 185, 1989). The DNA primer to be used for the amplification is one which is complementary to the 3' ends of the double stranded DNA containing an entire region of the *lysC* gene or a segment thereof. If only a portion of the *lysC* gene is to be amplified, then the gene library must be screened for DNA fragments containing the entire region, using the DNA fragment as the primer. If the entire region is to be amplified, then the DNA fragment may be subjected to agarose gel electrophoresis, and then the desired band is cut out to recover the DNA fragment containing the *lysC* gene.

The DNA primer may be appropriately prepared based on, for example, a known sequence of *E. coli* - (Cassan, M., Parsot, C., Cohen, G.N. and Patte, J.C., *J. Biol. Chem.*, 261, 1052, 1986), and preferred are the two types of primers, 5'-CTTCCCTTGTGCCAAGGCTG-3' (SEQ ID NO: 2) and 5'-GAATTCCTTTGCGAGCAG-3' (SEQ ID NO: 3), which are capable of amplifying the 1347-base region coding for the *lysC* gene. The synthesis of the DNA may be carried out in a conventional manner, using a DNA synthesizer Model 380B, produced by Applied Biosystems Co., and the phosphoramidite method (see Tetrahedron Letters 22, 1859, 1981). The PCR reaction may be carried out using a DNA thermal cycler Model PJ2000, produced by Takara Shuzo Co., and Taq DNA polymerase, produced by Takara Shuzo Co., following the method indicated by the supplier.

The *lysC* gene which has been amplified by the PCR method is linked to vector DNA which is capable of replication in *Escherichia* bacteria, and it is introduced into cells thereof. The method of transforming a host with the vector DNA to be used and the method of confirming the presence of *lysC* are the same as those described previously.

The method of inducing mutations into the obtained *lysC*, such as the substitution, insertion or deletion of amino acids, may be effected by the recombinant PCR method (Higuchi, R., 61, in PCR Technology (Erich, H.A., Eds., Stockton Press, 1989)), the site-directed mutagenesis method (Kramer, W. and Frits, H.J., Meth. in Enzymol., 154, 350, 1987; Kunkel, T.A. et al., Meth. in Enzymol., 154, 367, 1987), etc. By using these methods, a desired mutation may be induced at a desired site. Furthermore, for the introduction of random mutation may be used a method of directly treating the object gene on chromosome DNA or a plasmid with hydroxylamine (Hashimoto, T. and Sekiguchi, M., J. Bacteriol., 159, 1039, 1984); a conventional method involving exposure of cells containing the object DNA to ultraviolet rays or to treatment with a chemical agent such as N-methyl-N'-nitrosoguanidine, nitrous acid, or the like; or a method of chemically synthesizing the object gene.

The method of selecting the inhibition-releasing mutant gene *lysC*^{*} involves, at first, transforming with the mutated recombinant DNA an AK-deficient strain, for example, *E. coli* GT3 strain. Next, the transformant is cultured in a minimal medium, for example, M9, containing a significant amount of lysine. The strains which possess plasmids having the wild type of *lysC* exhibit the inhibition by lysine of AK III which is the only AK in the strain, and therefore the synthesis of threonine, isoleucine, methionine and diaminopimelic acid (DAP) is no longer possible and growth is inhibited. In contrast, the strains which possess plasmids having *lysC*^{*} which releases the inhibition by lysine should be capable of growing on a minimal medium containing a significant amount of lysine. Utilizing this phenomenon, the desired strains, that is, strains with plasmids having the *lysC*^{*} which releases the inhibition, i.e., those which are resistant to lysine or to S-2-aminoethylcysteine (AEC), an analogue of lysine, may be selected.

The above mentioned mutant gene obtained in this manner may be used as the recombinant DNA for introduction into an appropriate microorganism (host) and expressed therein to obtain a microorganism possessing AK with release of the feedback inhibition.

The above mentioned wild strains of *Escherichia* bacteria may be mentioned as hosts into which the obtained *lysC* gene or mutant *lysC* gene (*lysC*^{*}) is to be introduced and then amplified for the production of threonine, but in addition to these, any other bacteria may be used as hosts provided that both the replication origin of the recombinant vector DNA constructed here and the *lysC* gene or the mutated *lysC* gene (*lysC*^{*}) function therein, the recombinant vector DNA is capable of replication therein, and the *lysC* gene or the mutated *lysC* gene (*lysC*^{*}) may be reinforced therein. The most preferable host is *E. coli* B-3996 strain.

The transformants obtained in the above manner, which possess the recombinant vector DNA containing the gene which codes for aspartokinase III with release of the feedback inhibition due to lysine, are cultured, and the object L-threonine is produced and accumulated in the culture solution and then collected.

The culture medium to be used for the production of L-threonine is a conventional culture medium containing a carbon source, a nitrogen source, inorganic ions, and if necessary other organic components.

The carbon source to be used may be a saccharide such as glucose, lactose, galactose, fructose, a hydrolyzed starch, an alcohol such as glycerol or sorbitol, or an organic acid such as fumaric acid, citric acid, succinic acid, or the like.

The nitrogen source to be used may be an inorganic ammonium salt such as ammonium sulfate, ammonium chloride, ammonium phosphate, etc., an organic nitrogen such as soybean hydrolysate, or ammonia gas, ammonia water, or the like.

A required substance such as vitamin B1, L-homoserine or the like, or yeast extract, is preferably added in an appropriate amount as an organic trace nutrient source. In addition to these, potassium phosphate, magnesium sulfate, iron ion, manganese ion, etc. are added in small amounts if necessary.

The culturing is preferably carried out under aerobic conditions for 16-72 hours, while the culturing temperature is controlled to 25°C-45°C, and the pH of the culture to 5-8. For the control of the pH, an inorganic or organic acid, alkali substance, or ammonia gas, etc. may be used. The collection of the L-threonine from the fermented mash is carried out by combining the conventional ion exchange resin method, precipitation method and any other publicly known methods.

(Brief Description of the Drawings)

Fig. 1 is a schematic diagram of the biosynthetic pathway of threonine.

Fig. 2 is a restriction enzyme map for pLYSC1 and pLYSC2 (See Example 1).

Fig. 3 is a graph showing the effects of the introduction of hydroxylamine-induced mutations on *lysC* (See Example 2 (2-4)).

Fig. 4 shows the degree of lysine-dependent inhibition of aspartokinases coded for by various *lysC*^{*} genes (See Example 3 (3-4)). The lysine concentrations on the horizontal axis are expressed as logarithms.

The specific activities on the vertical axis are expressed as ratios defining as 100% the activity of a wild type of AK III with 0 mM addition of lysine.

Fig. 5 shows a diagram for the construction of pLLC* (See Example 4 (4-1)).

Fig. 6 shows a diagram for the construction of pVICLC*A and pVICLC*B (See Example 4 (4-3)).

5 Fig. 7 shows the points of mutation of each LysC* (See Example 5 (5-3)).

(Best Mode for Carrying Out the Invention)

A more concrete description of the present invention is provided below, with reference to the Examples.

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Example 1: Cloning of the wild lysC gene

The base sequence of the lysC gene of *E. coli* has already been reported (Cassan, M., Parsot, C., Cohen, G.N. and Patte, J.C., J. Biol. Chem., 261, 1052, 1986), and it is known that its open reading frame (ORF) has 1,347 bases coding for 449 amino acids. Since an operator is present which is responsible for repression by lysine, this operator region was removed, and the cloning was effected by amplifying only the region containing the SD sequence and the ORF, using the PCR method.

Two different primers, 5'-CTTCCCTTGTCGAAGGCTG-3' (Sequence No. 2) and 5'-GAATTCCTTTGC-GAGCAG-3' (Sequence No. 3) were prepared, and the entire genomic DNA of *E. coli* K-12 MC1061 strain was recovered using the method of Saito and Miura (Biochem. Biophys. Acta., 72, 619, 1963). These were used in a PCR reaction according to the method of Erlich, et al. (PCR Technology, Stockton Press, 1989) to amplify the object DNA. The resulting DNA was digested with *Bam*HI and *Ase*I, and then the ends thereof were made blunt and the DNA was inserted into the *Sma*I site of the multicopy vector pUC18. Thus were obtained two types of plasmids, one antisense (pLYSC1) and one sense (pLYSC2) with respect to the lacZ promoter (Fig. 2).

25 These plasmids were used for transformation of *E. coli* GT3 strains (thrA1016^b, metLM1005, lysC1004) completely deficient in AK I, II and III, and since the homoserine + diaminopimeric acid requirements of GT3 were complemented, this confirmed that the gene was lysC which codes for the active AK III.

30 Example 2: Acquisition of inhibition-released mutant gene lysC*

(2-1);

To efficiently obtain an inhibition-released lysC gene (lysC*), the gene on the recombinant plasmid DNA prepared in Example 1 was subjected to mutating treatment.

35 As a method of obtaining the inhibition-released mutant gene lysC*, first the wild lysC recombinant plasmid was incorporated for transformation into the completely AK-deficient *E. coli* GT3. A significant amount of lysine was added to an M9 minimal medium having the composition listed in Table 1 below, and the transformants were cultured in the medium. The strains possessing plasmids having the wild type of lysC exhibited inhibition of the only AK, i.e., AK III, by lysine, and therefore the synthesis of threonine, isoleucine, methionine and diaminopimeric acid (DAP) was no longer possible and growth was inhibited.

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Table 1: M9 minimal medium

5	A:20 x M9	(g/l)
	Na ₂ HPO ₄ · 12H ₂ O	303
10	KH ₂ PO ₄	60
	NaCl	10
	NH ₄ Cl	20
15	B:1 M MgSO ₄	
	C:50% glucose	
20	D:1 g/l thiamine	

25 A, B, C, D and water were sterilized separately and
 mixed at a ratio of A:B:C:D:water = 5:0.1:1:0.1:95.

30 In contrast, the strains which possess plasmids having lysC⁺ which release the inhibition by lysine are expected to be capable of growth in a minimal medium containing a significant amount of lysine. Utilizing this phenomenon, selection was made of strains whose growth was resistant to lysine or to S-2-aminoethylcysteine (AEC), an analogue of lysine, i.e., those strains with plasmids having the lysC⁺ which releases the inhibition.

35 (2-2);Investigation of the conditions of selection of the feedback inhibition-resistant lysC mutants (lysC⁺)

40 First, pLYSC1 and pLYSC2 were each incorporated into *E. coli* GT3 for its transformation to obtain two different transformants, and culturing was effected on an M9 minimal agar plate medium containing lysine or AEC. Also, the growth inhibition concentration of lysine or AEC was determined, and the conditions for selection of lysC⁺ were investigated.

45 As shown in Table 2, it was clear that with respect to pLYSC2 the amount of expression was amplified by the lacZ promoter, and thus there was resistance to a considerably high concentration of lysine or AEC even with the wild type of lysC, but that since the lysC gene of pLYSC1 was antisense with respect to the lacZ promoter and the promoter of the LysC gene itself was lacking, the amount of expression was low and the growth was inhibited under a lower concentration of lysine or AEC (growth was completely inhibited by addition of either lysine or AEC at about 0.2 mM. The "+" in the table indicates growth of the transformant, and the "-" indicates no growth thereof). This growth inhibition was found to be restored by the simultaneous addition of homoserine and DAP.

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Table 2: Relationship between growth of completely AK-deficient GT3 strain possessing lysC plasmid and concentration of added lysine or lysine analogue AEC

(a): Lys concentration and growth

	0	0.2	0.4	0.8	1.5	3	6	12	25	50	100	200 (mM)
GT3/pLYSC1	+	-	-	-	-	-	-	-	-	-	-	-
GT3/pLYSC2	+	+	+	+	+	+	+	+	+	+	±	-

(b): AEC concentration and growth

	0	0.2	0.4	0.8	1.5	3	6	12	25	50 (mM)
GT3/pLYSC1	+	-	-	-	-	-	-	-	-	-
GT3/pLYSC2	+	±	±	±	±	±	-	-	-	-

M9 agar medium, 37°C, 2 day culturing

+: growth, ±: some growth, -: no growth

Thus, pLYSC1 was used for the mutation introduction experiment, and the selection medium used for selection of the lysC⁺ was prepared by adding 10 mM of lysine or 0.2 mM of AEC to the M9 minimal medium.

(2-3); Mutating treatment

Hydroxylamine is a chemical mutating agent which induces a mutation of C→T by converting cytosine to N⁴-hydroxycytosine. For the introduction of the mutation into the plasmid, two methods were used, one an *in vitro* mutating treatment method whereby the plasmid was treated directly with hydroxylamine, and the other an *in vivo* mutating treatment method to provide variety to the mutation, i.e., for mutations other than C→T, whereby cells possessing the plasmids are treated with nitrosoguanidine (NTG) and then the plasmids are extracted.

(2-4); *In vitro* mutating treatment with hydroxylamine

A 2 µg portion of the DNA was treated in a reaction solution shown in Table 3 below, i.e., in 0.4 M of hydroxylamine at 75°C for 1-4 hours. The treated DNA was purified with glass powder, and then used for the transformation into a completely AK-deficient GT3 strain, which was plated onto a complete medium (L-broth; 1% bacto trypton, 0.5% yeast extract, 0.5% NaCl, 1.5% agar) to form colonies. The formed colonies were replicated onto the selection medium obtained in (2-1). The ratio of appearance of the transformants and the mutation ratio varied as shown in Fig. 3. With 4 hours of treatment, the yield of the mutant strain was 0.5-0.8%, which is a rather high value.

Table 3

Reaction solution composition	
0.1 M KH ₂ PO ₄ - 1 mM EDTA (pH 6.0)	100 µl
1 M hydroxylamine - 1 mM EDTA (pH 6.0)	80 µl
DNA	2 µg
Water	balance
Total	200 µl

(2-5); *In vivo* mutating treatment with nitrosoguanidine (NTG)

pLYSC1 was used for transformation of *E. coli* MC1061, and the cells were directly subjected to treatment with NTG. The treated cells were cultured overnight for fixation of the mutation, after which the plasmids were extracted therefrom, and used for transformation into GT3, and screening was carried out in the same manner as in (2-4) to obtain lysine resistant (Lys^R) or AEC resistant (AEC^R) mutants. The treatment is outlined in Table 4 below.

Table 4: Outline of treatment

(1) Culturing of cells containing the object plasmid in a 2 x TY medium (1.6% bacto trypton, 1% yeast extract, 0.5% NaCl), followed by cell collection at O.D. 660 nm = approx. 0.3.

(2) Washing with TM buffer.

(3) Suspension in NTG solution (0.2 mg/ml in TM buffer), followed by treatment at 37°C for 0-90 minutes.

(4) Washing with TM buffer and 2 x TY medium, followed by culturing overnight in 2 x TY medium and fixation of mutation.

(5) Extraction of plasmid DNA from cells, transformation of the GT3 strain, and screening for recombinant strains.

The composition of the TM buffer was as follows;

Tris	50 mM
Maleic acid	50 mM
(NH ₄) ₂ SO ₄	1 g/l
MgSO ₄ • 7H ₂ O	0.1 g/l
Ca(NO ₃) ₂	5 mg/l
FeSO ₄ • 7H ₂ O	0.25 mg/l
Adjusted to pH 6.0 using NaOH.	

Example 3: Isolation of lysC⁺ gene

(3-1);

A total of 180 candidate strains obtained in Example 2 (hydroxylamine-treated = 48 strains, and NTG-treated = 132 strains) were again spotted onto a selection medium, and the presence of AEC or lysine resistance was confirmed to obtain 153 strains. Taking note of the difference in the patterns of amino acid accumulation in the medium, the 153 strains were separated into 14 groups and a representative of each group was selected for the measurement of AK activity. Since there was no great difference between the mutants treated with hydroxylamine and those treated with NTG, the experiment was continued without distinction between the two.

(3-2); Measurement of aspartokinase activity

A completely AK-deficient GT3 strain was used as the host in respect of which the above mentioned 14 mutant pLYSC1's (named the pLYSC1⁺ series) and wild pLYSC1 plasmids were respectively used for transformation, after which a cell-free extract was prepared from the transformants and the enzyme activity of AK III was measured. The method for the measurement of the enzyme activity was as follows.

(3-3); Measurement method for AK III activity

(3-3-1); Method for preparation of crude enzyme solution

- (1) Culturing of cells in 2 x TY medium, followed by cell collection at O.D. 660 nm = approx. 0.8.
- (2) Washing with 0.02 M KH₂PO₄ (pH 6.75) - 0.03 M β mercaptoethanol at 0 °C.
- (3) Sonic crushing of cells (0 °C, 100 W, 30 sec x 4 times).
- (4) Centrifugation at 0 °C, 33 krpm for 1 hour, followed by addition of ammonium sulfate to supernatant to 80% saturation.
- (5) Centrifugation, followed by dissolution of pellets in buffer of (2), and preservation (-20 °C).

(3-3-2); Method for measurement of enzyme activity

The method of measuring the enzyme activity was according to the method of Stadtman, et al. (Stadtman, E.R., Cohen, G.N., Lebras, G. and Robichon-Szulmajster, H., J. Biol. Chem., 236, 2033, 1961).

The reaction solution had the composition shown in Table 5 below.

Table 5

Reaction solution composition	
Reaction mixture* ¹	0.3 ml
Hydroxylamine solution* ²	0.2 ml
0.1 M potassium aspartate (pH 7.0)	0.1 ml
Enzyme solution	
Water	balance
Total	1.0 ml
The reaction solution without potassium aspartate is defined as blank.	

*1: 9 ml of 1 M Tris-HCl (pH 8.1) + 0.5 ml of 0.3 M MgSO₄ + 5 ml of 0.2 M ATP (pH 7.0)

*2: 8 M hydroxylamine neutralized with KOH just prior to use

The measurement method is outlined in Table 6 below.

Table 6: Outline of measurement method

(1) Incubation of the reaction solution at 27°C for 45 minutes.

(2) Addition of an FeCl₃ solution (0.4 ml of 2.8 N HCl

+ 0.4 ml of 12% TCA + 0.7 ml of 5% FeCl₃·6H₂O/0.1 N HCl) for coloration.

(3) Centrifugation, followed by measurement of absorbance value at 540 nm (A_{540}) of supernatant.

The activity was expressed as the amount of hydroxamic acid produced during one minute.

1 U = 1 μ mol/min. Molar absorbance coefficient = 600.

The results are shown in Fig. 4.

(3-4); Degree of release of lysine-dependent inhibition

For the measurement of the enzyme activity of AK, lysine was added to the enzyme reaction solutions at various concentrations, and the levels of lysine-dependent inhibition were determined. The wild type of AK III was very strongly inhibited by lysine, e.g., with a 50% inhibition with about 0.45 mM of lysine, and

roughly 100% inhibition with 5 mM thereof (Fig. 4).

In contrast, the mutants of AK III obtained here exhibited various degrees of release, but all 14 had some release of the lysine-dependent inhibition (Fig. 4, and Table 7). Particularly, in the case of Nos. 24, 80, 117, 169 and 172, practically no inhibition was observed even with 100 mM of lysine, and the 50% inhibition concentration was 200 fold or greater.

Also, the specific activity, the activity of the enzyme per unit protein, was equal to or greater than that of the wild type, even considering the influence of the growing conditions of the cells and the preparation of the sample, and there was seen no problem of reduced activity due to introduction of the mutations (Table 7). From this it was assumed that the active site of AK III and the site responsible for regulation by lysine are independent from each other.

In Table 7, the inhibition release is presented as the activity (%) in the presence of 100 mM of lysine with respect to the activity of a lysine-free reaction solution, and the thermal stability is presented as the maintenance of activity (%) after treatment at 55 °C for 1.5 hours with respect to the activity without heating.

Table 7

Inhibition release by lysine and thermal stability			
	Specific activity (μ /mg protein)	Inhibition release (%)	Thermal stability (%)
Wild type	0.0247	0	18
No. 117	0.0069	120	0
No. 24	0.0218	100	30
No. 80	0.0244	99	36
No. 172	0.0189	97	0
No. 169	0.0128	96	2
No. 150	0.0062	77	25
No. 126	0.0250	61	39
No. 149	0.0256	59	9
No. 167	0.0083	43	45
No. 48	0.0228	38	42
No. 60	0.0144	35	9
No. 158	0.0224	22	42
No. 156	0.0101	18	2
No. 43	0.0212	17	0

(3-5); Thermal stability

When raising the level of activity of certain enzymes for their improvement, it is important that they be stably maintained within the cells. Because of the difference in the activity of proteases within and without the cells and the influence of the preservation buffer, measurement thereof is preferably made *in vivo*, but here for the sake of convenience, thermal stability as a parameter of each of the mutants of AK III was tested *in vitro*.

As a result of various tests regarding the temperature at which the inactivation of AK III occurs, the temperature was set to 56 °C and the rate of maintenance of activity was determined after the treatment effected for 90 minutes. As shown in Table 7 above, half of the mutants were superior to the wild type. In general, mutated proteins tend to be less stable than their wild types, but the some of the mutated types of AK III obtained here had a greater stability than the wild type, and many were thought to be very useful as enzymes for the practical production of L-threonine.

Example 4: Fermentation production of L-threonine using a lys⁺C-introduced strain

(4-1);

Of the heretofore known threonine-producing *E. coli* bacteria, the B-3996 strain exhibits the highest capability of threonine production. Here, it was decided to use the B-3996 strain as the host for the evaluation of lys⁺C. The B-3996 strain has been deposited at the Research Institute of Genetics and

Selection of Industrial Microorganism under No. БКПМ (VKPM) B-3996. Furthermore, as the *lysC*^{*} to be evaluated were provided 6 types with different degrees of inhibition release and specific activities (Nos. 24, 43, 60, 80, 149 and 167 in Table 7) for use in the following experiment.

First, in order to increase the degree of expression of the *lysC*^{*}, each of the above mentioned 6 types of *lysC*^{*} on pLYSC1^{*} was transferred to the downstream of the lacZ promoter of the vector pHSG399 (product of Takara Shuzo Co.) to effect the inverting insertion of each *lysC*^{*}. The novel plasmids obtained in this manner were collectively named pLLC^{*} series (Fig. 5). The *E. coli* HB101 strains into which the plasmids having the above *lysC*^{*} Nos. 80 and 167 had been inserted were named AJ12750 and AJ12751, respectively, and they were originally deposited at the National Institute of Bioscience and Human Technology (Japan) as of September 1, 1992. The AJ12750 was assigned No. FERM P-13136 (converted to an international deposit as of November 4, 1993 under No. FERM BP-4462), and the AJ12751 was assigned No. FERM 13137 (converted to an international deposit as of the same date under No. FERM BP-4463). The rest were not deposited, but since all of the mutation points for each *lysC*^{*} have been discovered, as described below, those having ordinary skill in the art may easily recover the plasmids from the above mentioned deposited bacteria following the method of Maniatis, et al. (Sambrook, J., Fritsch, E.F., Maniatis, T., Molecular Cloning, Cold Spring Harbor Laboratory Press, 1, 21, 1989), to obtain the object rest *lysC*^{*} gene following the site directed mutagenesis method (Sambrook, J., Fritsch, E.F., Maniatis, T., Molecular Cloning, Cold Spring Harbor Laboratory Press, 15, 83, 1989). A conventional method was used to insert these plasmids into the B-3996 strain for evaluation.

The culturing was carried out using the following culture medium as shown in Table 8 below (the components A, B and C having been separately sterilized), for a culturing time of 38 hours, at a temperature of 37 °C with stirring at 114-116 rpm.

Table 8

25

30

35

40

Threonine production medium		
		(g/l)
A:	(NH ₄) ₂ SO ₄	16
	KH ₂ PO ₄	1
	MgSO ₄ · 7H ₂ O	1
	FeSO ₄ · 7H ₂ O	0.01
	MnSO ₄ · 5H ₂ O	0.01
	Yeast ext. (Difco)	2
	L-Met	0.5
Adjusted to pH 7.0 with KOH, and autoclaved at 115 °C for 10 minutes (16/20 vol).		
B:	20% glucose, autoclaved at 115 °C for 10 minutes (4/20 vol).	
C:	Pharmacopoeial CaCO ₃ , autoclaved at 180 °C for 2 days (30 g/l).	
D:	Antibiotics (100 µg/ml of streptomycin, and 5 µg/ml of kanamycin).	

45 (4-2); Evaluation of *lysC*^{*}

Each of the six pLLC^{*} series plasmids was used for the transformation of the B-3996 strain, and each transformant was cultured under both conditions of addition of 1 g/l of lysine and no addition thereof, respectively. As controls, the host B-3996 strains with and without a plasmid possessing the wild type of *lysC* (pLLC1) were prepared.

The results are shown in Table 9 below. In the table, the values in the parentheses are the ratios of lysine yields (g) per 1g consumed saccharide to the lysine yield (g) per 1g consumed saccharide obtained when the B-3996 was cultured with no addition of lysine (control). This means that the latter yield is referred to as 1.00. The lysine sensitivity is defined as (lysine yield per consumed saccharide with addition of lysine)/(lysine yield per consumed saccharide with no addition of lysine). With respect to the B-3996 strain, the reduction in the yield per consumed saccharide observed in the lysine-added culture was approximately 0.74 on the basis of that observed in the non-added culture. However, in the case of the strains into which a *lysC*^{*} had been introduced, for example, B-3996/pLLC^{*}149, the reduction in the yield per consumed

saccharide observed in the lysine-added culture was approximately 0.96 on the basis of that observed in the non-added culture. From this it is supposed that the AK III encoded by *lysC* contributes to the biosynthesis of threonine, and that the inhibition on AK III activity by the addition of lysine is, in turn, related to the inhibition on threonine biosynthesis. This phenomenon is diminished when the *lysC*^{*} according to the present invention is used.

Table 9

	Amount of Lys added (g/l)	Residual saccharide (g/l)	Thr (g/l)	Yield per consumed saccharide (%)	Lysine sensitivity
B-3996	0	0.09	12.1	30.7 (1.00)	
	1	0.12	8.9	22.6	0.74
B-3996/pLLC1 (wild)	0	0.07	13.1	33.0 (1.07)	
	1	0.14	11.1	28.3	0.87
B-3996/pLLC [*] 24	0	0.11	15.4	38.5 (1.25)	
	1	0.08	12.7	32.3	0.84
B-3996/pLLC [*] 43	0	0.09	15.1	38.3 (1.25)	
	1	0.12	14.4	35.5	0.93
B-3996/pLLC [*] 60	0	1.32	13.8	36.3 (1.18)	
	1	2.66	11.6	31.4	0.86
B-3996/pLLC [*] 80	0	0.10	15.7	39.7 (1.29)	
	1	0.09	13.9	35.6	0.90
B-3996/pLLC [*] 149	0	0.07	12.8	32.1 (1.04)	
	1	1.80	11.3	30.7	0.96
B-3996/pLLC [*] 167	0	0.07	15.5	39.8 (1.30)	
	1	0.10	13.9	35.3	0.89

Also, regarding the threonine production of the lysine-free culture, it was clear that the strains which had the wild *lysC* plasmid gave improved productivity over the B-3996 strain having no such plasmids, and that the productivity was further increased by the strains having mutant *lysC*^{*}s (approximately 1.3 times that of the host in case of No. 80). These results indicate that AK III is one of the rate-limiting factors in the production of threonine by the B-3996 strain. With the cells into which the mutant *lysC*^{*} had been introduced, the lysine-free culturing gave an improved yield compared to that using the cells into which the wild *lysC* had been introduced, and it is thought that this was due to the fact that the latter exhibited inhibition on the wild type of AK III by lysine biosynthesized in the cells themselves.

(4-3); Stabilization of mutant *lysC*^{*} plasmids

Plasmids as shown in Fig. 6 were prepared with *lysC*^{*} incorporated in both directions at the *Bam*HI site of the plasmid pVIC40 extracted from the B-3996 strain. The method for collecting pVIC40 from the B-3996 strain was already described above. Selection was made of No. 80, which had a high level of threonine production, as the *lysC*^{*} and of No. 43 for comparison therewith, to obtain 4 different plasmids, i.e., pVICLC^{*}80A, pVICLC^{*}80B, pVICLC^{*}43A and pVICLC^{*}43B.

For use as the host, B-399 strain (B-3996 strain with pVIC40 removed) was newly prepared by curing the B-3996 strain. The curing was achieved by repeating a 1/100-fold dilution three times with a streptomycin-free L-broth, with respect to a culture solution of the B-3996 strain. The culturing temperature here was 40°C. Into the B-399 strain obtained in this manner were introduced pVICLC^{*}80A and pVICLC^{*}43A, respectively, out of the 4 plasmids having *lysC*^{*} incorporated therein, and culturing was effected. The results are shown in Table 10. The strains into which had been introduced pVICLC^{*}80A or

pVICLC⁺43A exhibited an increase in yield over the pVIC40 transformant strain (B-3996) being used as the control (1.10-fold with respect to pVICLC⁺80A, and 1.07-fold with respect to pVICLC⁺43A).

Table 10

Plasmid	Residual saccharide (g/l)	Thr (g/l)	Yield per consumed saccharide (%)
pVIC40	0.15	14.0	35.7 (1.00)
pVICLC ⁺ 43A	0.09	14.9	38.0 (1.07)
pVICLC ⁺ 80A	0.09	15.3	39.1 (1.10)
pVIC40 + pLLC ⁺ 43	0.97	15.8	41.1 (1.15)
pVIC40 + pLLC ⁺ 80	6.15	13.7	41.3 (1.16)

It is thought that the reduction in the degree of increase in the yields, compared with the results in (4-2), was due to a lower number of copies resulting from changing of the vector to pVIC40, but the growth was the same as for the B-3996 strain and the plasmids were stably maintained.

(4-4); Incorporation of mutant lysC⁺ gene into chromosomes

The mutant type lysC⁺ was introduced, utilizing the homologous recombination phenomenon, into the threonine-producing B-3996 by targeting the wild lysC gene on its chromosome. The method used here was a modification of the method of Russel et al. (Russel, M. and Model, P., J. Bacteriol., 159, 1034, 1984).

If the host is an *E. coli* mutant, i.e., the MM382 strain (po1A^{ts}, available from the *E. coli* genetic stock center, U.S.A.), then the plasmid pHSG399 can replicate at 37 °C (permissive temperature), but not at high temperatures such as 42 °C (non-permissive temperature). This fact was utilized for the homologous recombination.

First, pLLC43⁺ and pLLC80⁺ were each introduced into the mutant strain MM383 as the host, at a temperature of 37 °C at which replication was possible, to obtain transformants MM383/pLLC⁺ 43 and MM383/pLLC⁺ 80, respectively. These transformants were cultured at 42 °C, and selection was made of those which had lacked the plasmids but still remained resistant to chloramphenicol. The plasmids had been incorporated into the chromosomes of these strains. The obtained strains were named MM383-C43 and MM383-C80, respectively. These strains were infected with P1 phage, and a phage solution was prepared, which was used to infect the B-3996 strain. Selection was made of strains which had lysC⁺ transduced into their chromosomes, using chloramphenicol resistance as the marker, and they were named B-3996-C43 and B-3996-C80, respectively.

Culturing of B-3996-C-43 and B-3996-C-80 was effected under the same conditions as in (4-1), and the results shown in Table 11 below were obtained. Approximately a 4% improvement in the yield of threonine production was observed.

Table 11

	Amount of Lys added (g/l)	Thr (g/l)	Yield per consumed saccharide (%)
B-3996	0	14.1	34.8 (1.00)
	1	9.9	24.6
B-3996-C43	0	14.3	35.4 (1.02)
	1	11.4	28.1
B-3996-C80	0	14.6	36.1 (1.04)
	1	11.2	27.7

Example 5: Determination of base sequences of wild *lysC* and mutant *lysC**

(5-1);

5 Using a DNA sequencer "ABI Model 373A" (ex ABI Co.), a conventional method was conducted to determine the base sequence of the wild *lysC* gene. The result is listed in the attached sequence listing under SEQ ID NO: 1. It was found that the sequence are different from the already published base sequence of *lysC* of *E. coli* K-12 JC411 (Cassan, M., Parsot, C., Cohen, G.N. and Patta, J.C., J. Biol. Chem., 261, 1052, 1986) in 6 bases (resulting in two amino acid residue changes). The 6 differences were
10 thought attributable to the difference in the strains used.

(5-2); Base sequence of the feedback inhibition-resistant mutant *lysC**

15 The base sequences of the 14 kinds of *lysC** investigated in Example 3 were determined in the same manner as in (5-1), and the points of mutation were clearly found. The results are shown in Table 12. Of the 14 kinds, two exactly identical pairs were present, and thus there were 12 kinds of actual mutants. Mutant Nos. 149, 150, 156, 158, 167, 169 and 172 had been obtained by treatment with hydroxylamine and Nos. 24, 43, 48, 60, 80, 117 and 126 by treatment with NTG. However, the patterns of mutation were all C→T or G→A; the mutation of G→A had been given through the mutation of C→T on the complimentary strand.

Table 12

Identification of Mutation Points of <i>lysC</i> *		
Mutant types <i>lysC</i> *	Mutagen (a)	Mutation point (amino acid change)
No. 126	N	GGT→GA [*] T (³²³ Gly→Asp)
No. 43	N	GGT→GA [*] T (³²³ Gly→Asp) GGC→GA [*] C (⁴⁰⁸ Gly→Asp)
No. 149	H	CGT→T [*] GT (³⁴ Arg→Cys) GGT→GA [*] T (³²³ Gly→Asp)
No.43/167 NO. 150	N/H H	CTC→T [*] TC (³²⁵ Leu→Phe) ATG→ATA [*] (³¹⁸ Met→Ile)
No. 172	H	⁷⁷⁵ C→T (silent) ATG→ATA [*] (³¹⁸ Met→Ile) GTG→A [*] TG (³⁴⁹ Val→Met)
No. 117 No. 158 No. 24/80	N H N/N	TCA→TT [*] A (³⁴⁵ Ser→Leu) GTG→A [*] TG (³⁴⁷ Val→Met) ACC→AT [*] C (³⁵² Thr→Ile)
No. 169	H	⁹²³ C→T (silent) ACC→AT [*] C (³⁵² Thr→Ile) TCT→TT [*] T (³⁵⁹ Ser→Phe)
No. 60	N	⁸⁵⁹ G→A (silent) GAA→A [*] AA (¹⁵⁴ Glu→Lys)
No. 156	H	ATG→ATA [*] (⁴¹⁷ Met→Ile) TGT→TA [*] T (⁴¹⁹ Cys→Tyr) ²⁰¹⁴ C→T (silent)

(a) H: hydroxylamine treatment, N: NTG treatment

55 (5-3); Identification of domains contributing to release of feedback inhibition

A large number of mutations were present at the C-end, and they were particularly concentrated at the A domain (from the 318th Met residue to the 325th Leu residue) and the B domain (from the 345th Ser

residue to the 352nd Thr residue) of Fig. 7 (see "Mutation points" column in Table 12). *E. coli* has two types of AK, trhA (AK I-HD I) and metL(M) (AK II-HD II), in addition to lysC (AK III). The mutant lysC* had two locations having a high degree of homology to the two types, and these domains are thought to be the active center of AK (Cassan, M., Parsot, C., Cohen, G.N. and Patte, J.C., J. Biol. Chem., 261, 1052, 1986).

5 The A and B domains at the C-end are outside of these common domains, and thus it is highly possible that they are lysine-controlled domains specific to AK III. Particularly, Nos. 24/80, 172, 169 and 117 with a high degree of inhibition release had mutations in the B domain, which fact is thought to be important. Of the 12 types, 10 had mutations in either the A or B domain. Nos. 60 and 158 were exceptional, and showed low levels of both release of inhibition and thermal stability (Table 7 above), and this is thought to be a
10 result of partial release of the inhibition due to a change in the structure of the enzyme as a whole by mutation. Also, in the case of the mutant types having multiple mutation points, there is the question of which mutation point is effective, and the following explanation may be given based on comparison with the mutant types having a single mutation at the same site.

15 (1) Nos. 126 (1 point), 43 (2 points) and 149 (2 points)

The mutation point of No. 126 in the A domain is common to all the three mutant types, while the degree of inhibition release is the same for Nos. 149 and 126, and the greater number of mutation points in No. 43 resulted in rather opposite effect. This leads to the assumption that the mutation point of No. 126 in
20 the A domain is the one which imparts the inhibition release.

(2) Nos. 24/80 (1 point) and 169 (2 points)

Of the two mutation points in No. 169, the one in the B domain was the same as that in No. 24/80. Both
25 produced a 100% degree of inhibition release, and therefore the mutation point in No. 24/80 is sufficient.

(3) Nos. 150 (1 point) and 172 (2 points)

No. 150 has one mutation point in the A domain, while No. 172 has two mutations each in the A domain
30 (same as No. 150) and the B domain, respectively. The degree of inhibition release was stronger with No. 172, making it clear that both the A domain and the B domain contribute to the inhibition release.

In the manner described above, the present inventors have identified the domain relating to the release of the lysine-dependent inhibition of aspartokinase activity in the A domain (when expressed in terms of amino acid residues, from the 318th Met residue to the 325th Leu residue) and the B domain (from the
35 345th Ser residue to the 352nd Thr residue). However, the present inventors have not concluded that the release of the lysine-dependent inhibition of aspartokinase activity occurs only in the case of mutation wherein some specific amino acid residues are changed to some other amino acid residues. For example, in Table 12 it is shown that release of the feedback inhibition occurs when the 323rd glycine residue is changed to an aspartic acid residue, but it is clear to a person having ordinary skill in the art that the same
40 release occurs even when it is replaced by a glutamic acid residue instead of an aspartic acid residue. This is because plural amino acids which have a similar structure, such as aspartic acid and glutamic acid, are called homologous amino acids, and exchange of an amino acid for its homologous one is not considered to cause any great change in the function of the protein (A list of homologous amino acids is found in Protein Engineering, p.31, CMC Co., 1985).

45 It is often observed that exchange of an amino acid for a non-homologous amino acid has the same effect as that for a homologous one. Conversely, exchange of an amino acid for a homologous amino acid sometimes produces dramatic changes in the function of the protein (Estell, D.A., Graycar, T.P., and Wells, J.A., J. Biol. Chem., 260, 6518, 1988; Schultz, S.C., and Richards, J.H., Procedure. Natl. Acad. Sci. USA, 83, 1588, 1986; Yutani, K., et al., J. Biol. Chem., 262, 13429, 1987; Yutani, K., et al., Procedure. Natl. Acad.
50 Sci. USA, 84, 4441, 1987; Nishiyama, M., et al., J. Biol. Chem., 266, 14294, 1991).

Furthermore, according to the present invention, some of the amino acid residues in the A and B domains were not changed, but since the functions of the proteins are defined on the basis of domain units, it is reasonable to assume that release of feedback inhibition will be observed by introducing one or more amino acid residue mutations in the domains which have not been introduced this time. In fact, a number of
55 such examples have been reported (Furuya, H., et al., Biochemistry, 28, 6848, 1989; Furuya, H., et al. Biochem. Biophys. Research. Commun., 160, 669, 1989; Cunningham, B.C. and Wells, J.A., Science, 244, 1081, 1989).

In short, the present invention is characteristically based on the present inventor's discovery of the domains relating to the feedback inhibition. In the Examples, only a few patterns of mutation in the domains are disclosed, but as described above, it is clear to a person having ordinary skill in the art that other patterns of mutation may produce the same effect. Therefore, such mutations are also within the scope of the present invention.

(Industrial Applicability)

As mentioned above, *Escherichia* bacteria-derived AK III genes have been obtained which sufficiently release the feedback inhibition due to lysine. By introducing these genes into threonine-producing bacteria, it is possible to obtain other threonine-producing bacteria which were much improved in threonine production over those of the prior art. The successful use of these threonine-producing bacteria has provided a much more excellent method for the production of L-threonine by fermentation than the conventional methods.

Sequence Listing

SEQ ID NO:1

Sequence length:2147

Sequence Type:nucleic acid

Strandedness:double

Topology:linear

Molecule Type:Genomic DNA

Original Source:

Organism:Escherichia coli

Strain:MC1061

Feature:

Feature Key:-35 signal

Position:242..249

Feature Determination Method: S

Feature:

Feature Key:-10 signal

Position:265..273

Feature Determination Method: S

Feature:

Feature Key:primer bind

Position:536..555

Feature Determination Method: E

Feature:

Feature Key:primer bind

Position:2128..2147

Feature Determination Method: E

Feature:

Feature Key:RBS

Position:575..578

Feature Determination Method: S

Feature:

Feature Key:mat peptide

Position:584..1930

Feature Determination Method: S

Feature:

Feature Key:terminator

Position:1941..1968

Feature Determination Method: S

Sequence Description

TCGAAGTGTT TCTGTAGTGC CTGCCAGGCA GCGGTCTGCG TTGGATTGAT GTTTTTCATT	60
AGCAATACTC TTCTGATTTT GAGAATTGTG ACTTTGGAAG ATTGTAGCGC CAGTCACAGA	120
AAAATGTGAT GGTTTTAGTG CCGTTAGCGT AATGTTGAGT GTAAACCCTT AGCGCAGTGA	180
AGCATTTATT AGCTGAACTA CTGACCGCCA GGAGTGGATG AAAAATCCGC ATGACCCCAT	240

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	CGTTGACAAC CGCCCCGCTC ACCCTTTATT TATAAATGTA CTACCTGCGC TAGCGCAGGC	300
	CAGAAGAGGC GCGTTGCCCA AGTAACGGTG TTGGAGGAGC CAGTCCTGTG ATAACACCTG	360
5	AGGGGGTGCA TCGCCGAGGT GATTGAACGG CTGGCCACGT TCATCATCGG CTAAGGGGGC	420
	TGAATCCCCT GGGTTGTCAC CAGAAGCGTT CGCAGTCGGG CGTTTCGCAA GTGGTGGAGC	480
	ACTTCTGGGT GAAAATAGTA GCGAAGTATC GCTCTGCGCC CACCCGTCTT CCGCTCTTCC	540
10	CTTGTGCCAA GGCTGAAAAT GGATCCCCTG ACACGAGGTA GTT ATG TCT GAA ATT	595
	Met Ser Glu Ile	
	1	
	GTT GTC TCC AAA TTT GGC GGT ACC AGC GTA GCT GAT TTT GAC GCC ATG	643
15	Val Val Ser Lys Phe Gly Gly Thr Ser Val Ala Asp Phe Asp Ala Met	
	5 10 15 20	
	AAC CGC AGC GCT GAT ATT GTG CTT TCT GAT GCC AAC GTG CGT TTA GTT	691
	Asn Arg Ser Ala Asp Ile Val Leu Ser Asp Ala Asn Val Arg Leu Val	
	25 30 35	
20	GTC CTC TCG GCT TCT GCT GGT ATC ACT AAT CTG CTG GTC GCT TTA GCT	739
	Val Leu Ser Ala Ser Ala Gly Ile Thr Asn Leu Leu Val Ala Leu Ala	
	40 45 50	
	GAA GGA CTG GAA CCT GGC GAG CGA TTC GAA AAA CTC GAC GCT ATC CGC	787
25	Glu Gly Leu Glu Pro Gly Glu Arg Phe Glu Lys Leu Asp Ala Ile Arg	
	55 60 65	
	AAC ATC CAG TTT GCC ATT CTG GAA CGT CTG CGT TAC CCG AAC GTT ATC	835
	Asn Ile Gln Phe Ala Ile Leu Glu Arg Leu Arg Tyr Pro Asn Val Ile	
	70 75 80	
30	CGT GAA GAG ATT GAA CGT CTG CTG GAG AAC ATT ACT GTT CTG GCA GAA	883
	Arg Glu Glu Ile Glu Arg Leu Leu Glu Asn Ile Thr Val Leu Ala Glu	
	85 90 95 100	
	GCG GCG GCG CTG GCA ACG TCT CCG GCG CTG ACA GAT GAG CTG GTC AGC	931
35	Ala Ala Ala Leu Ala Thr Ser Pro Ala Leu Thr Asp Glu Leu Val Ser	
	105 110 115	
	CAC GGC GAG CTG ATG TCG ACC CTG CTG TTT GTT GAG ATC CTG CGC GAA	979
	His Gly Glu Leu Met Ser Thr Leu Leu Phe Val Glu Ile Leu Arg Glu	
	120 125 130	
40	CGC GAT GTT CAG GCA CAG TGG TTT GAT GTA CGT AAA GTG ATG CGT ACC	1027
	Arg Asp Val Gln Ala Gln Trp Phe Asp Val Arg Lys Val Met Arg Thr	
	135 140 145	

	AAC	GAC	CGA	TTT	GGT	CGT	GCA	GAG	CCA	GAT	ATA	GCC	GCG	CTG	GCG	GAA	1075
	Asn	Asp	Arg	Phe	Gly	Arg	Ala	Glu	Pro	Asp	Ile	Ala	Ala	Leu	Ala	Glu	
	150						155					160					
5	CTG	GCC	GCG	CTG	CAG	CTG	CTC	CCA	CGT	CTC	AAT	GAA	GCG	TTA	GTG	ATC	1123
	Leu	Ala	Ala	Leu	Gln	Leu	Leu	Pro	Arg	Leu	Asn	Glu	Gly	Leu	Val	Ile	
	165					170					175					180	
	ACC	CAG	GGA	TTT	ATC	GGT	AGC	GAA	AAT	AAA	GGT	CGT	ACA	ACG	ACG	CTT	1171
10	Thr	Gln	Gly	Phe	Ile	Gly	Ser	Glu	Asn	Lys	Gly	Arg	Thr	Thr	Thr	Leu	
					185					190						195	
	GGC	CGT	GGA	GGC	AGC	GAT	TAT	ACG	GCA	GCC	TTG	CTG	GCG	GAG	GCT	TTA	1219
	Gly	Arg	Gly	Gly	Ser	Asp	Tyr	Thr	Ala	Ala	Leu	Leu	Ala	Glu	Ala	Leu	
				200					205					210			
15	CAC	GCA	TCT	CGT	GTT	GAT	ATC	TGG	ACC	GAC	GTC	CCG	GGC	ATC	TAC	ACC	1267
	His	Ala	Ser	Arg	Val	Asp	Ile	Trp	Thr	Asp	Val	Pro	Gly	Ile	Tyr	Thr	
			215					220					225				
	ACC	GAT	CCA	CGC	GTA	GTT	TCC	GCA	GCA	AAA	CGC	ATT	GAT	GAA	ATC	GCG	1315
20	Thr	Asp	Pro	Arg	Val	Val	Ser	Ala	Ala	Lys	Arg	Ile	Asp	Glu	Ile	Ala	
		230					235					240					
	TTT	GCC	GAA	GCG	GCA	GAG	ATG	GCA	ACT	TTT	GGT	GCA	AAA	GTA	CTG	CAT	1363
	Phe	Ala	Glu	Ala	Ala	Glu	Met	Ala	Thr	Phe	Gly	Ala	Lys	Val	Leu	His	
						250					255					260	
25	CCG	GCA	ACG	TTG	CTA	CCC	GCA	GTA	CGC	AGC	GAT	ATC	CCG	GTC	TTT	GTC	1411
	Pro	Ala	Thr	Leu	Leu	Pro	Ala	Val	Arg	Ser	Asp	Ile	Pro	Val	Phe	Val	
					265					270					275		
	GGC	TCC	AGC	AAA	GAC	CCA	CGC	GCA	GGT	GGT	ACG	CTG	GTG	TGC	AAT	AAA	1459
30	Gly	Ser	Ser	Lys	Asp	Pro	Arg	Ala	Gly	Gly	Thr	Leu	Val	Cys	Asn	Lys	
				280					285					290			
	ACT	GAA	AAT	CCG	CCG	CTG	TTC	CGC	GCT	CTG	GCG	CTT	CGT	CGC	AAT	CAG	1507
	Thr	Glu	Asn	Pro	Pro	Leu	Phe	Arg	Ala	Leu	Ala	Leu	Arg	Arg	Asn	Gln	
			295					300					305				
35	ACT	CTG	CTC	ACT	TTG	CAC	AGC	CTG	AAT	ATG	CTG	CAT	TCT	CGC	GGT	TTC	1555
	Thr	Leu	Leu	Thr	Leu	His	Ser	Leu	Asn	Met	Leu	His	Ser	Arg	Gly	Phe	
			310				315					320					
	CTC	GCG	GAA	GTT	TTC	GGC	ATC	CTC	GCG	CGG	CAT	AAT	ATT	TCG	GTA	GAC	1603
40	Leu	Ala	Glu	Val	Phe	Gly	Ile	Leu	Ala	Arg	His	Asn	Ile	Ser	Val	Asp	
						330					335					340	

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	TTA ATC ACC ACG TCA GAA GTG AGC GTG GCA TTA ACC CTT GAT ACC ACC	1651
	Leu Ile Thr Thr Ser Glu Val Ser Val Ala Leu Thr Leu Asp Thr Thr	
	345 350 355	
5	GGT TCA ACC TCC ACT GGC GAT ACG TTG CTG ACG CAA TCT CTG CTG ATG	1699
	Gly Ser Thr Ser Thr Gly Asp Thr Leu Leu Thr Gln Ser Leu Leu Met	
	360 365 370	
	GAG CTT TCC GCA CTG TGT CGG GTG GAG GTG GAA GAA GGT CTG GCG CTG	1747
10	Glu Leu Ser Ala Leu Cys Arg Val Glu Val Glu Glu Gly Leu Ala Leu	
	375 380 385	
	GTC GCG TTG ATT GGC AAT GAC CTG TCA AAA GCC TGC GGC GTT GGC AAA	1795
	Val Ala Leu Ile Gly Asn Asp Leu Ser Lys Ala Cys Gly Val Gly Lys	
	390 395 400	
15	GAG GTA TTC GGC GTA CTG GAA CCG TTC AAC ATT CGC ATG ATT TGT TAT	1843
	Glu Val Phe Gly Val Leu Glu Pro Phe Asn Ile Arg Met Ile Cys Tyr	
	405 410 415 420	
	GGC GCA TCC AGC CAT AAC CTG TGC TTC CTG GTG CCC GGC GAA GAT GCC	1891
20	Gly Ala Ser Ser His Asn Leu Cys Phe Leu Val Pro Gly Glu Asp Ala	
	425 430 435	
	GAG CAG GTG GTG CAA AAA CTG CAT AGT AAT TTG TTT GAG TAAATACTGT	1940
	Glu Gln Val Val Gln Lys Leu His Ser Asn Leu Phe Glu	
	440 445	
25	ATGGCCTGGA AGCTATATTT CGGGCCGTAT TGATTTTCTT GTCACATATGC TCATCAATAA	2000
	ACGAGCCTGT ACTCTGTTAA CCAGCGTCTT TATCGGAGAA TAATTGCCTT TAATTTTTTTT	2060
	ATCTGCATCT CTAATTAATT ATCGAAAGAG ATAAATAGTT AAGAGAAGGC AAAATGAATA	2120
30	TTATCAGTTC TGCTCGCAAA GGAATTC	2147

SEQ ID NO:2

Sequence length:20

Sequence Type:nucleic acid

Strandedness:single

35 Topology:linear

Molecule Type:Other nucleic acid, synthetic DNA

Sequence Description

CTTCCCTTGT GCCAAGGCTG

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SEQ ID NO:3

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Sequence length:18

Sequence Type:nucleic acid

Strandedness:single

Topology:linear

Molecule Type:Other nucleic acid, synthetic DNA

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Sequence Description

GAATTCCTTT GCGAGCAG

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55 Claims

1. A DNA containing a gene which codes for aspartokinase III originating in *Escherichia* bacteria and which has in the coding domain a mutation which releases the feedback inhibition by lysine on said

aspartokinase III.

2. A DNA according to Claim 1, wherein the mutation which releases the feedback inhibition by lysine on aspartokinase III is positioned on the A domain or B domain of aspartokinase III.
- 5 3. A DNA according to Claim 1, wherein the mutation which releases the feedback inhibition by lysine on aspartokinase III is selected from the group consisting of a mutation wherein the 323rd Gly is replaced by Asp; a mutation wherein the 323rd Gly is replaced by Asp and the 408th Gly is replaced by Asp; a mutation wherein the 34th Arg is replaced by Cys and the 323rd Gly is replaced by Asp; a mutation wherein the 325th Leu is replaced by Phe; a mutation wherein the 318th Met is replaced by Ile; a mutation wherein the 318th Met is replaced by Ile and the 349th Val is replaced by Met; a mutation wherein the 345th Ser is replaced by Leu; a mutation wherein the 347th Val is replaced by Met; a mutation wherein the 352nd Thr is replaced by Ile; a mutation wherein the 352nd Thr is replaced by Ile and the 369th Ser is replaced by Phe; a mutation wherein the 164th Glu is replaced by Lys; and a mutation wherein the 417th Met is replaced by Ile and the 419th Cys is replaced by Tyr, in Sequence Listing: SEQ ID NO: 1.
- 10 4. A recombinant DNA wherein a DNA according to any of Claims 1, 2 and 3 is linked to a vector DNA capable of autonomous replication in an *Escherichia* bacterial cell.
- 20 5. A microorganism belonging to the genus *Escherichia* which has been transformed by the introduction into its cell of the recombinant DNA according to Claim 4.
- 25 6. A microorganism according to Claim 5 which has been transformed by the incorporation of a DNA according to any of Claims 1, 2 and 3 into its chromosomal DNA.
- 30 7. A method for the production of L-threonine which comprises culturing a microorganism according to Claim 5 or 6 in a fermentation medium, whereby L-threonine is produced and accumulated in the culture medium, and collecting the L-threonine from said culture medium.

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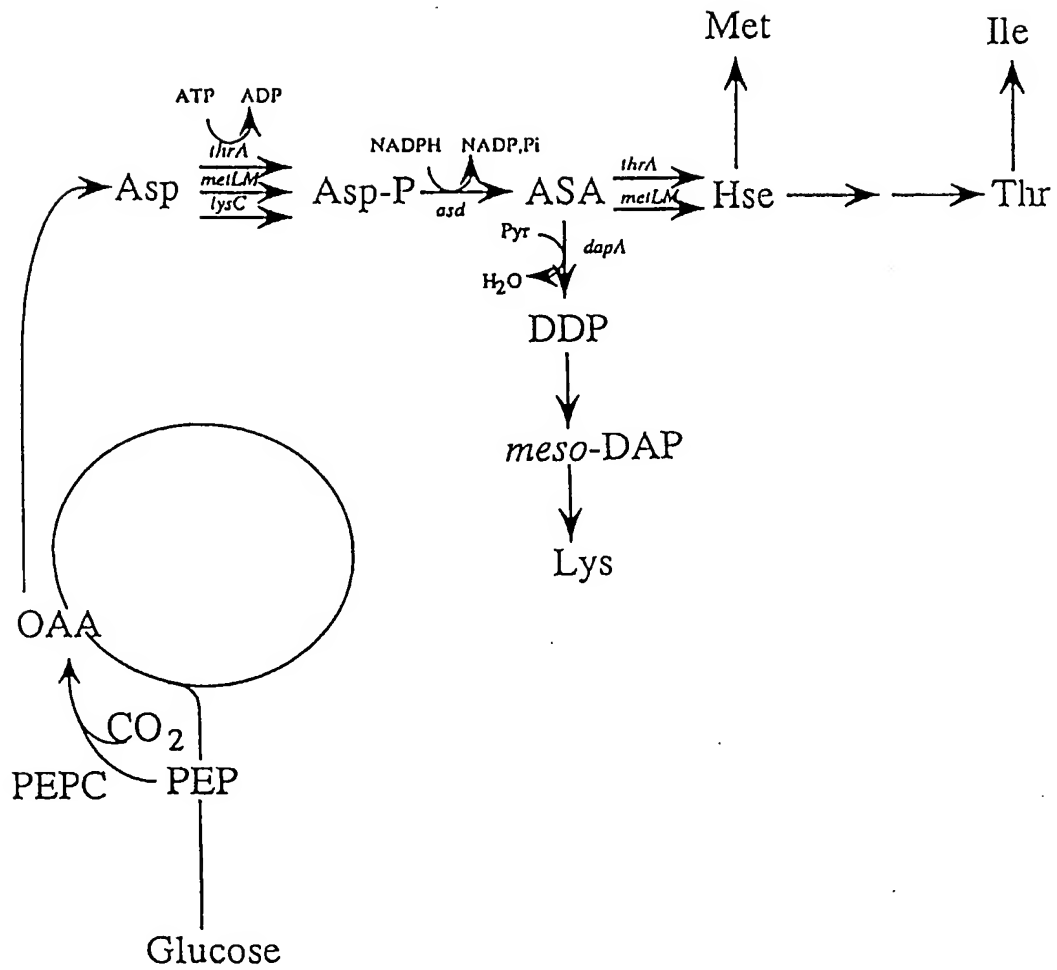


Fig.1

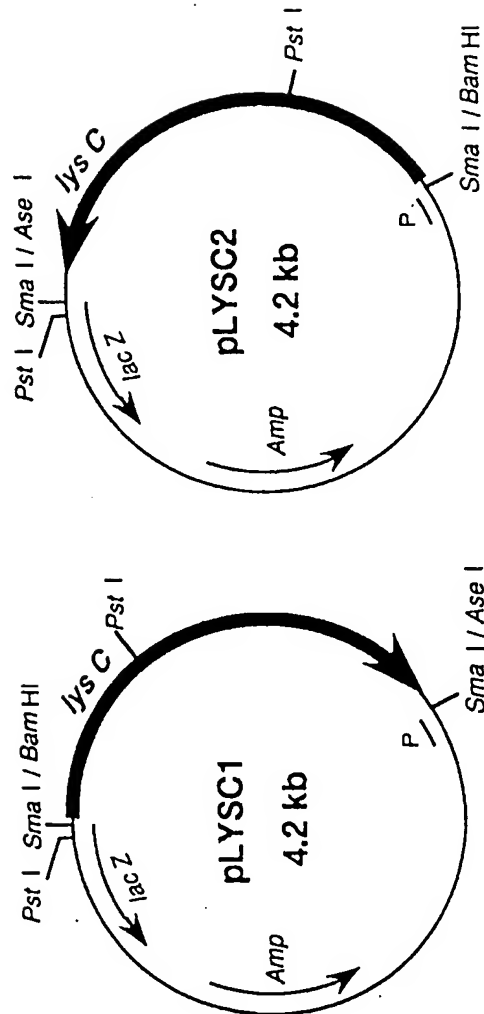


Fig. 2

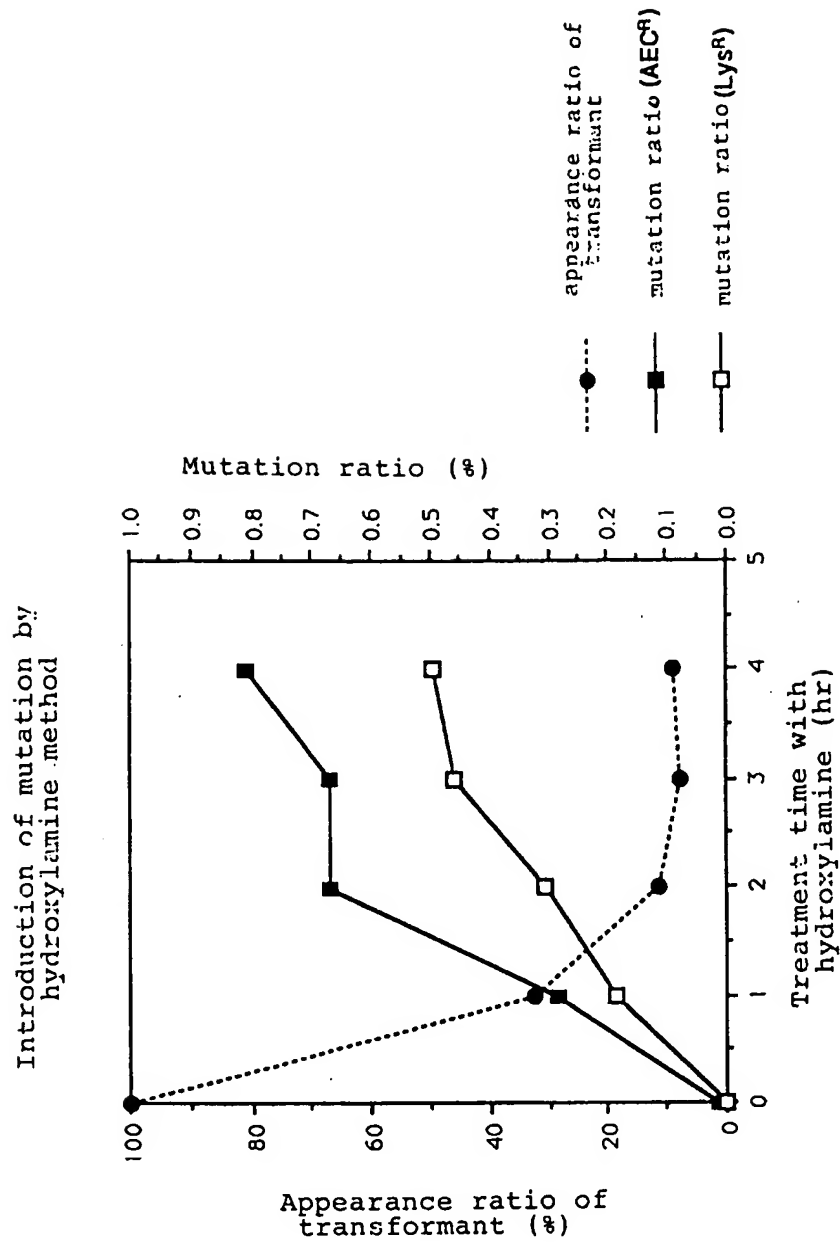
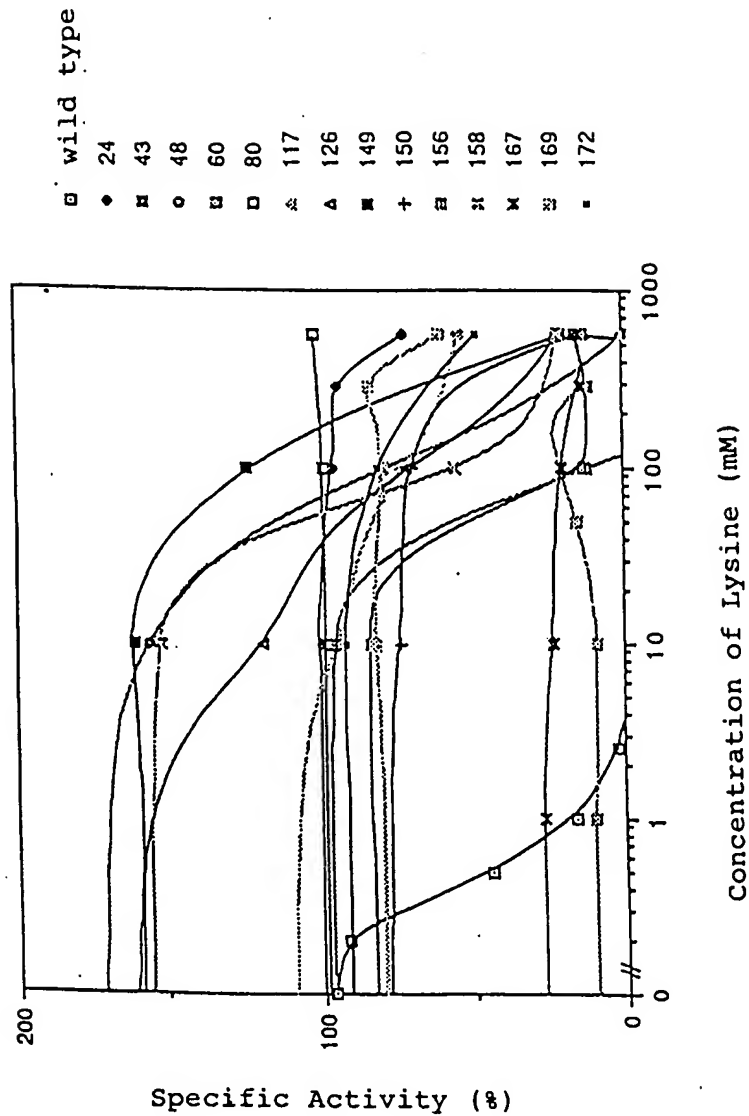


Fig. 3

Fig. 4



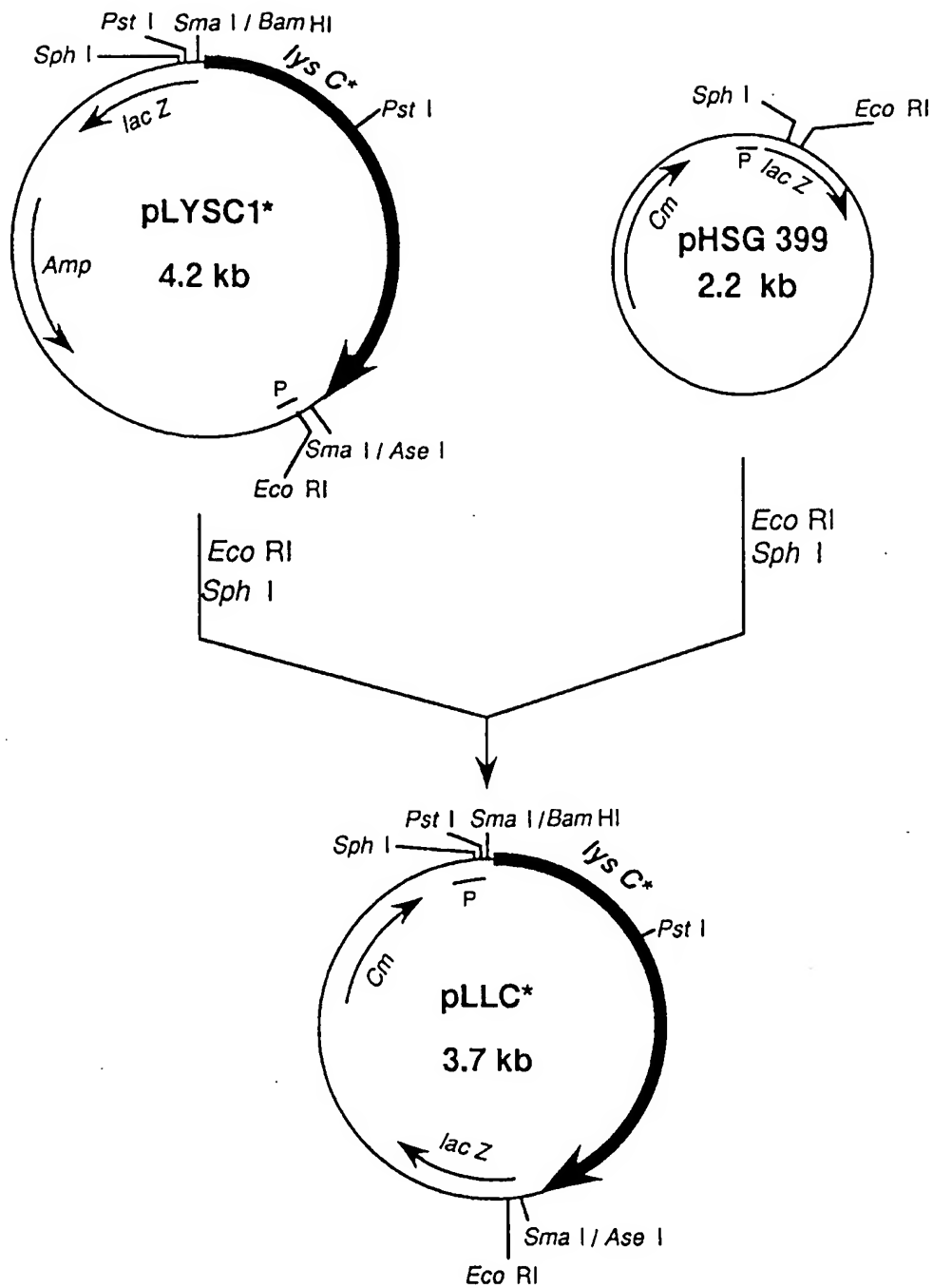


Fig. 5

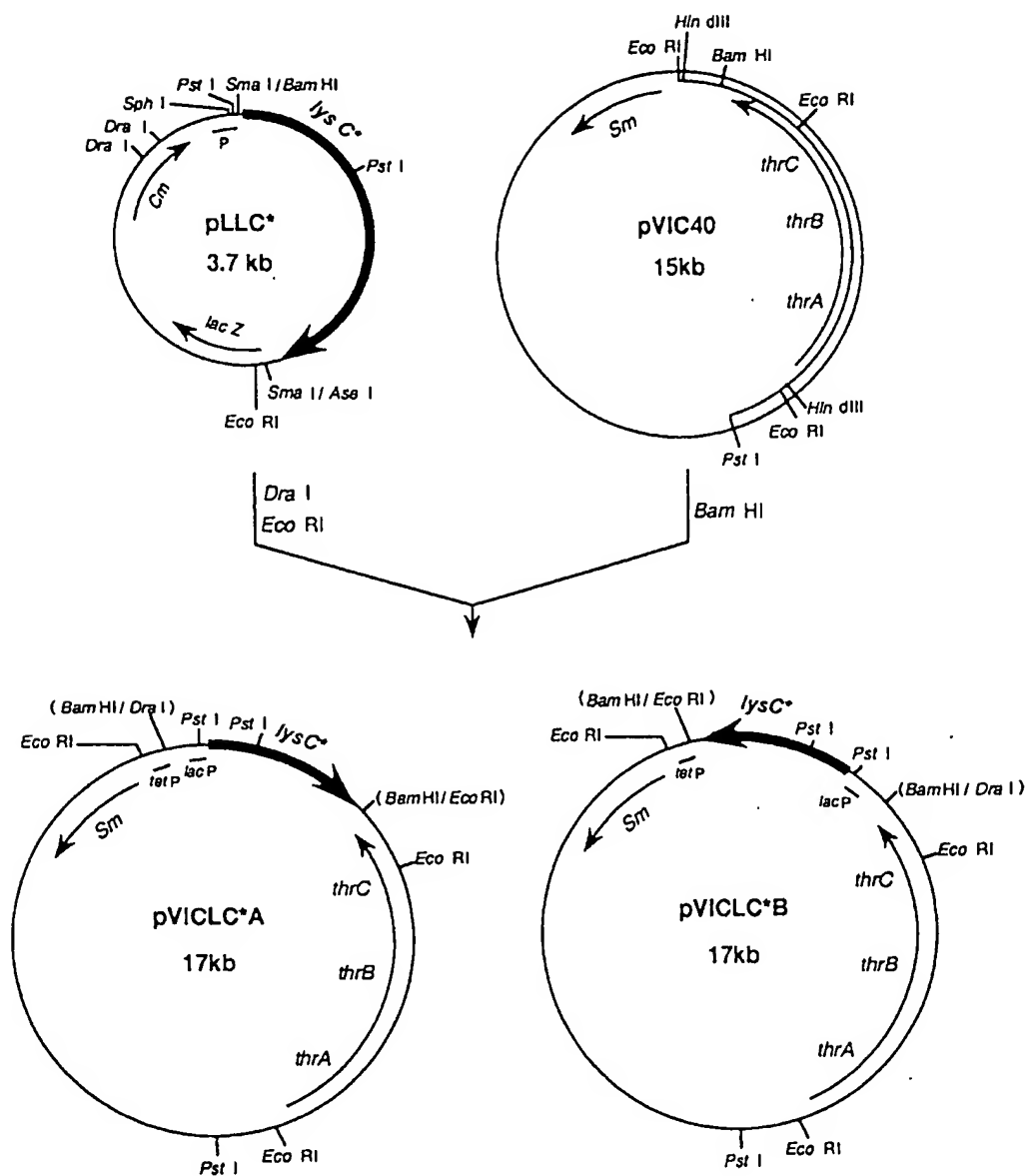


Fig. 6

AAT CCG CCG CTG TTC CGC GCT CTG GCG CTT CGT CGC AAT CAG ACT CTG	1513
Asn Pro Pro Leu Phe Arg Ala Leu Ala Leu Arg Arg Asn Gln Thr Leu	
300 310	
<div style="text-align: center;"> A A T ↑ ↑ ↑ (A) </div>	
CTC ACT TTG CAC AGC CTG AAT <u>ATG CTG CAT TCT CGC GGT TTC CTC</u> GCG	1561
Leu Thr Leu His Ser Leu Asn Met Leu His Ser Arg Gly Phe Leu Ala	
↓ ↓ ↓	
Ile Asp Phe	
320	
GAA GTT TTC GGC ATC CTC GCG CGG CAT AAT ATT TCG GTA GAC TTA ATC	1609
Glu Val Phe Gly Ile Leu Ala Arg His Asn Ile Ser Val Asp Leu Ile	
330 340	
<div style="text-align: center;"> T A A T ↑ ↑ ↑ ↑ (B) </div>	
ACC ACG <u>TCA GAA GTG AGC GTG GCA TTA ACC</u> CTT GAT ACC ACC GGT TCA	1657
Thr Thr Ser Glu Val Ser Val Ala Leu Thr Leu Asp Thr Thr Gly Ser	
↓ ↓ ↓ ↓	
Leu Met Met Ile	
350	
ACC TCC ACT GGC GAT ACG TTG CTG ACG CAA TCT CTG CTG ATG GAG CTT	1750
Thr Ser Thr Gly Asp Thr Leu Leu Thr Gln Ser Leu Leu Met Glu Leu	
360 370	

Fig. 7

INTERNATIONAL SEARCH REPORT

International application No.

PCT/JP93/01640

A. CLASSIFICATION OF SUBJECT MATTER Int. Cl ⁵ C12N15/54, C12N1/21, C12P13/08// (C12N15/54, C12R1:19), (C12N1/21, C12R1:19), (C12P13/08, C12R1:19) According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) Int. Cl ⁵ C12N15/54, C12N1/21, C12P13/08 Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) BIOSIS PRE VIEWS, CAS ONLINE		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	THE Journal of Biological Chemistry, Volume 261, No. 3, (Jan. 25, 1986), Michel Cassan et al., "Nucleotide Sequence of lysC gene encoding the Lysine-sensitive Aspartokinase III of Escherichia coli K12", see p. 1052-1057	1-5
A	JP, B1, 1-29559 (Ajinomoto Co., Inc.), June 12, 1989 (12. 06. 89), & FR, A, 2453216 & US, A, 4347318 & GB, B, 2049670 & DE, C, 3012921	5, 7
A	Agricultural and Biological Chemistry, Volume 50, No. 4 (April, 1986), Tamio Mizukami et al. "Essential role of Aspartokinase in L-Threonine production by Escherichia coli W mutants", see p. 1015-1018	1-7
<input type="checkbox"/> Further documents are listed in the continuation of Box C. <input type="checkbox"/> See patent family annex.		
* Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family		
Date of the actual completion of the international search		Date of mailing of the international search report
January 27, 1994 (27. 01. 94)		February 15, 1994 (15. 02. 94)
Name and mailing address of the ISA/ Japanese Patent Office		Authorized officer
Facsimile No.		Telephone No.

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